


```

FT      sig-peptide      /product- "Human full-length 36PIG3/SGP28 protein"
FT      3...98      /*tag- b
FT      mat-peptide      99...776
FT      /*tag- c
FT      /product- "Human mature full-length 36PIG3/SGP28 protein"
XX      WO200131343-A2.
XX      03-MAY-2001.
XX      27-OCT-2000: 2000WO-US293607.
XX      28-OCT-1999: 99US-0162610.
XX      (UROC-) UROGENESYS INC.
XX      Hubert RS, Raitano AB, Afar DEH, Mitchell SC, Faris M;
XX      Jakobovits A.
XX      WPI: 2001-308685/32.
XX      P-PSDB: AAE02211.
PT      Detecting cancers, particularly of prostate and colon, from
PT      overexpression of SGP28 protein, also methods for treating these
PT      cancers e.g. by vaccination with the protein
XX      Claim 16: Page 62-63; 102pp; English.
XX      The present invention relates to methods and compositions for the
XX      diagnosis and therapy of prostate cancer which utilize human SGP28
XX      (specific granule protein 28) gene and proteins. The method involves
XX      detecting cancers, particularly of prostate and colon, from
XX      overexpression of SGP28 protein. The expression of SGP28, which is an
XX      extracellular protein is restricted to the prostate and ovary, and is
XX      markedly up-regulated in prostate tumours. SGP28 sequence is used for
XX      diagnosis (including in vivo imaging), staging, monitoring and prognosis
XX      of prostatic and colon cancer, and for assisting selection of therapy.
XX      Also SGP28-expressing cancers can be treated by administering a
XX      composition or vaccine that contains a vector expressing an antibody
XX      specific for SGP28 protein, nucleic acid encoding SGP28 protein or its
XX      fragments, polypeptides encoded by SGP28 gene and SGP28-specific antibody
XX      optionally conjugated to toxin or therapeutic agent. SGP28 gene product
XX      is also used as source of therapeutic antisense or ribozyme agents, as
XX      primers/probes for diagnosis or prognosis, to identify compounds that
XX      inhibit calcium entry into prostatic cells, for recombinant production
XX      of SGP28 peptides and for isolating related sequences. SGP28 protein and
XX      its fragments are used to raise specific antibodies (Ab) and to identify
XX      specific binding agents (potentially useful as therapeutic and
XX      diagnostic agents) and also potential anticancer agents. The present
XX      sequence is human full-length 36PIG3/SGP28 cDNA.
XX      SQ      Sequence 2144 BP; 735 A; 403 C; 382 G; 624 T; 0 other;
XX
XX      Alignment Scores:
XX      Pred. No.:      1.8e-136      Length:      2144
XX      Score:      1436.00      Matches:      258
XX      Percent Similarity:      100.00%      Conservative:      0
XX      Best Local Similarity:      100.00%      Mismatches:      0
XX      Query Match:      100.00%      Indels:      0
XX      DB:      22      Gaps:      0
XX
XX      US-09-698-781-3 (1-258) x AAD06222 (1-2144)
XX
XX      QY      1 MetLysGlnIleuNHISProAlaLeuGluThrThraIamethrLeuPheProValIleu 20
XX      |||||||
XX      DB      3 ATGAACAACAACTTCACTCCCTGGAACACACTGCAATGATATTTCCCACTGCTG 62
XX      |||||||
XX      QY      21 LeuPheLeuValAlaGlyLeuLeuProSerPheProAlaAsnGluAspLysAspProAla 40
XX      |||||||
XX      DB      63 TTCTTCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 122
XX      |||||||
XX      QY      41 PheThrIleLeuLeuLeuThrThrGlnThrGlnValGlnArgGluIleValAsnLysHisAsn 60
XX      |||||||
XX
XX      123 TTTACTGCTTTGTTTACCACCCCAACACAACTGCMAAGGGGATGTGTAATACCACT 182
XX      |||||||
XX      QY      61 GluLeuArgArgAlaValAsnSerProProAlaArgAsnMetLeuLysMetGluThrAsnLys 80
XX      |||||||
XX      DB      183 GAACTGAGAGAGACAGATATCTCCCTGCGCAAAACATGCTGAAAGATGGAATGGACAA 242
XX      |||||||
XX      QY      81 GluAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnTYrArgHisSerAsnPro 100
XX      |||||||
XX      DB      243 GAGCTGACGACAAATGCGCAAAAGCTGGGCAAAACAGTGCATTAACAGACAGTAACCA 302
XX      |||||||
XX      QY      101 LysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTYrMetSerSerAlaProSer 120
XX      |||||||
XX      DB      303 AAGGATGAATGACAGCTTMAATGTGTGAGATCTCTCAAGTCAAGTCCGCCAGC 362
XX      |||||||
XX      QY      121 SerTrpSerGlnAlaIleGlnSerTrpPheAspGluTYrAsnAspPheAspPheGlyVal 140
XX      |||||||
XX      DB      363 TCATGGTCACAGCAATCCAAAGCTGGTTGATGATGATGATGATGATGATGATGATGATG 422
XX      |||||||
XX      QY      141 GlyProLysThrProAsnAlaValAlaGlyHisTYrThrGlnValAlaTrpTYrSerSer 160
XX      |||||||
XX      DB      423 GGGCCAAAGACTCCCAACGCGAGTGTGACATTAACAGAGTGTGTGTGTGTGTGTGTGT 482
XX      |||||||
XX      QY      161 TyrLeuValAlaGlyCysGlyAsnAlaTYrCysProAsnGlnLysValLeuLysTYrTYr 180
XX      |||||||
XX      DB      483 TACCTCGTGGATGTGMAATGCTTACTGTCCTCAATCAAAAGTTCMAAATACTACTAT 542
XX      |||||||
XX      QY      181 ValCysGlnTYrCysProAlaGlyAsnTrpAlaAsnArgLeuTYrValProTYrGluGln 200
XX      |||||||
XX      DB      543 GTTGCCAAATATGCTCTGCTGTAATGGCTAATAGCTATATGTCCTTAAGAACAA 602
XX      |||||||
XX      QY      201 GlyAlaProCysAlaSerCysProAsnAsnCysAspAspGlyLeuCysThrAsnGlyCys 220
XX      |||||||
XX      DB      603 GGACACCTTGTGCGACAGTCCAGATTAACGTGACGATGATGATGATGATGATGATGATG 662
XX      |||||||
XX      QY      221 LysTYrGluAspLeuTYrSerAsnCysLysSerLeuLysLeuThrLeuThCysLysHis 240
XX      |||||||
XX      DB      663 AAGTACGAGAGATCTCTATAGTACTGTAAAGTTTGAAGTCAACATTAACCTGTAAACAT 722
XX      |||||||
XX      QY      241 GlnLeuValArgAspSerCysLysAlaSerCysAsnCysSerAsnSerIleTYr 258
XX      |||||||
XX      DB      723 CAGTTGGACAGGACAGTGTGCAAGCATCTCGCAATGTCTTAACACACATTAT 776
XX      |||||||
XX
XX      RESULT 2
XX      ABV24823
XX      ID      ABV24823 standard; cDNA; 1610 BP.
XX      XX
XX      AC      ABV24823:
XX      XX
XX      DT      16-SEP-2002 (first entry)
XX      XX
XX      DE      Human prostate expression marker cDNA 24814.
XX      XX
XX      KW      Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
XX      KM      pharmacogenomic marker; gene; ss.
XX      XX
XX      OS      Homo sapiens.
XX      XX
XX      PN      WO200160860-A2.
XX      PD      23-AUG-2001.
XX      PE      20-FEB-2001: 2001WO-US05171.
XX      PF      17-FEB-2000: 2000US-183319P.
XX      PR      16-MAR-2000: 2000US-189862P.
XX      PR      25-MAY-2000: 2000US-207454P.
XX      PR      09-JUN-2000: 2000US-211314P.
XX      PR      18-JUL-2000: 2000US-219007P.
XX      PR      13-DEC-2000: 2000US-255281P.
XX      PA      (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.

```


Best Local Similarity: 99.61% Mismatches: 1
 Query Match: 99.44% Indels: 0
 DB: 22 Gaps: 0

US-09-698-781-3 (1-258) x AAH98651 (1-2133)

```

Oy 1 MetysglnileuHnlsProAlaLeuGluThrAlaMetThrLeuPheProValleu 20
Db 2 ATGAACAATACTTCACTGCTGGAACCACTGCAATGACATTAATCCAGTCTG 61
Oy 21 LeupheleValAlaGlyLeuLeuProSerPheProAlaAsnGluAspLysAspProAla 40
Db 62 TTGTTCCGCTGCTGGGCTGCTTCATCTTTCCAGCAATGAAGATAGAGATCCCGCT 121
Oy 41 PheThraLeuLeuThrThrGlnThrGlnValGlnArgGluIleValAsnLysHisAsn 60
Db 122 TTTACTGCTTGTATACCAACCAACCAAGTGAAGGAGATGTGAATAGCACAAT 181
Oy 61 GluLeuArgArgAlaValSerProProAlaArgAsnMetLeuLysMetGluTrpAsnLys 80
Db 182 GAACGAGAGAGAGAGATCTCCCTCCAGAAACATGCGAATGGAATGGAACAA 241
Oy 81 GluAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnTyArgHisSerAsnPro 100
Db 242 GAGCGTCAGCAAAATGCCCAAAAGTGCGCAACAGTCATTAACAGACAGTAACCA 301
Oy 101 LysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTyMetSerSerAlaProSer 120
Db 302 AAGGATCAATAGCAAAAGCTAAATGTGTGAGATCTCTACATGTCAAGTCCCTCAGC 361
Oy 121 SerTrpSerGlnAlaIleGlnSerTrpPheAspGluTyArgAsnAspPheAspHelyal 140
Db 362 TCATGCTACAAAGCAATCCAAAGCTGTTGATGAGTACAAATGATTTGACTTGTGTA 421
Oy 141 GlyProLysThrProAsnAlaValAlaGlnHisTyThrGlnValAlaTrpTyrSerSer 160
Db 422 GGGCCAAAGATCCCAACAGCAGTGTGACATTATACAGAGTGTGTTGACTCTTCA 481
Oy 161 TyrLeuValAlaGlyCysGlyAsnAlaTyrcysProAsnGlnLysValLeuLysTyTrTy 180
Db 482 TACCTCGTGTGATGTGAAATGCCCTACCTGCCAATCAAAAAGTTCTAAATACACTAT 541
Oy 181 ValCysGlnTyrcysProAlaGlyAsnTrpAlaAsnArgLeuTyValProTyTrLueGln 200
Db 542 GTTGCCCAATATGCTCGCTGCTGATATGAGCTATATAGATGCTCCCTTATACAA 601
Oy 201 GlyAlaProCysAlaSerCysProAspAsnCysAspAspGlyLeuCysThraAsnGlyCys 220
Db 602 GGAGCAGCTTGTGCCAGTGTGCCAGATTAAGTGTGACATGACATGCAACCAATGCTGC 661
Oy 221 LysTrpGluAspLeuTyrcysAsnCysLysSerLeuLysLeuThrLeuThrcysLysHis 240
Db 662 AAGTACGAGAGATCTCTATAGTAACGTAAAGTTGAAGCTCACATTAACGCTAAACAT 721
Oy 241 GlnLeuValArgAspSerCysLysAlaSerCysAsnCysSerAsnSerIleTy 258
Db 722 CAGTTGGTCAGGAGACAGTTGCAAGGCAATCTGCAATTTGTTCAACAGCATTTAT 775

```

RESULT 4

AAH98659 standard; cDNA: 2133 BP.

AAH98659:

12-OCT-2001 (first entry)

Human EST-derived coding sequence SEQ ID NO: 516.

Human; sheep; pig; cow; fruit fly; yeast; hamster; macaque; horse;
 tomato; monkey; dog; sea urchin; expressed sequence tag; EST;
 diagnostics; forensic test; gene mapping; genetic disorder;
 biodiversity; gene therapy; nutrition; ss.

```

OS Homo sapiens.
XX WO200154477-A2.
XX 02-AUG-2001.
XX 25-JAN-2001; 2001WO-US02687.
XX 25-JAN-2000; 2000US-0491404.
XX 17-JUL-2000; 2000US-0617746.
XX 03-AUG-2000; 2000US-0631451.
XX 15-SEP-2000; 2000US-0663870.
XX (HXSE-) HXSEQ INC.
XX Tang YT, Liu C, Zhou P, Qian XB, Wang Z, Chen R, Asundi V;
XX Cao Y, Drmanac RA, Zhang J, Wehrman T;
XX WPI: 2001-476164/51.
XX P-PSDB: AAM24000.
XX Isolated polypeptide for treatment of diseases, diagnostics, raising
XX antibodies and research use -
XX Claim 1; Page 537-538; 1275pp; English.
XX The present invention provides the protein and coding sequences of novel
XX proteins from a variety of organisms, including human, dog, cat, horse,
XX cow, pig, hamster, monkey, macaque, yeast, bacteria, fruit fly, sea
XX urchin and tomato. These were derived from expressed sequence tags (ESTs)
XX from the organism of interest. They can be used in diagnostics,
XX forensics, gene mapping, identification of mutations, to assess
XX biodiversity and for nutritional purposes. The present sequence is a cDNA
XX of the invention.
XX SQ Sequence 2133 BP; 725 A; 403 C; 382 G; 623 T; 0 other;
XX
XX Alignment Scores:
XX Pred. No.: 1,17e-135 Length: 2133
XX Score: 1428.00 Matches: 257
XX Percent Similarity: 99.61% Conservative: 0
XX Best Local Similarity: 99.61% Mismatches: 1
XX Query Match: 99.44% Indels: 0
XX DB: 22 Gaps: 0
XX
XX US-09-698-781-3 (1-258) x AAH98659 (1-2133)
Oy 1 MetysglnileuHnlsProAlaLeuGluThrAlaMetThrLeuPheProValleu 20
Db 2 ATGAACAATACTTCACTGCTGGAACCACTGCAATGACATTAATCCAGTCTG 61
Oy 21 LeupheleValAlaGlyLeuLeuProSerPheProAlaAsnGluAspLysAspProAla 40
Db 62 TTGTTCCGCTGCTGGGCTGCTTCATCTTTCCAGCAATGAAGATAGAGATCCCGCT 121
Oy 41 PheThraLeuLeuThrThrGlnThrGlnValGlnArgGluIleValAsnLysHisAsn 60
Db 122 TTTACTGCTTGTATACCAACCAACCAAGTGAAGGAGATGTGAATAGCACAAT 181
Oy 61 GluLeuArgArgAlaValSerProProAlaArgAsnMetLeuLysMetGluTrpAsnLys 80
Db 182 GAACGAGAGAGAGAGATCTCCCTCCAGAAACATGCGAATGGAATGGAACAA 241
Oy 81 GluAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnTyArgHisSerAsnPro 100
Db 242 GAGCGTCAGCAAAATGCCCAAAAGTGCGCAACAGTCATTAACAGACAGTAACCA 301
Oy 101 LysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTyMetSerSerAlaProSer 120
Db 302 AAGGATCAATAGCAAAAGCTAAATGTGTGAGATCTCTACATGTCAAGTCCCTCAGC 361
Oy 121 SerTrpSerGlnAlaIleGlnSerTrpPheAspGluTyArgAsnAspPheAspHelyal 140

```


RESULT 6
 ABV24631
 ID ABV24631 standard; cDNA: 2452 BP.
 XX
 AC ABV24631;
 XX
 DT 16-SEP-2002 (first entry)
 XX
 DE Human prostate expression marker cDNA 24622.
 XX
 KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
 KW pharmacogenomic marker; gene; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200160860-A2.
 XX
 PD 23-AUG-2001.
 XX
 PF 20-FEB-2001; 2001WO-US05171.
 XX
 PR 17-FEB-2000; 2000US-183319P.
 PR 16-MAR-2000; 2000US-189862P.
 PR 25-MAY-2000; 2000US-207454P.
 PR 09-JUN-2000; 2000US-211314P.
 PR 18-JUL-2000; 2000US-219007P.
 PR 13-DEC-2000; 2000US-255281P.
 XX
 PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
 XX
 PI Schlegel R, Endege WO, Monahan JE;
 XX
 DR WPI: 2001-662795/76.
 XX
 PT Novel isolated nucleic acid molecule associated with cancerous state of
 PT prostate cells and correlating with presence of prostate cancer, useful
 PT for detecting presence of prostate cancer, stage of prostate cancer -
 XX
 PS Claim 1; Page 4680-4681; 11750pp; English.
 XX
 CC The invention relates to an isolated nucleic acid molecule (I) comprising
 CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
 CC specification or its complement. (I) is useful for:
 CC (a) assessing whether a patient is afflicted with prostate cancer;
 CC (b) monitoring the progression of prostate cancer in a patient;
 CC (c) assessing the efficacy of a test compound to inhibit prostate
 CC cancer in a patient;
 CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer
 CC in a patient;
 CC (e) selecting a composition for inhibiting prostate cancer in a patient;
 CC (f) assessing the prostate cell carcinogenic potential of a compound;
 CC (g) determining whether prostate cancer has metastasized in a patient;
 CC (h) assessing the aggressiveness or indolence of prostate cancer in a
 CC patient;
 CC (I) is also useful as a pharmacodynamic or pharmacogenomic marker.
 CC
 XX
 XX Sequence 2452 BP; 787 A; 467 C; 444 G; 754 T; 0 other;
 SQ
 Alignment Scores:
 Pred. No: 1.43e-135 Length: 2452
 Score: 1428.00 Matches: 257
 Percent Similarity: 99.61% Conservative: 0
 Best Local Similarity: 99.61% Mismatches: 1
 Query Match: 99.44% Indels: 0
 DB: 23 Gaps: 0
 US-09-698-781-3 (1-258) x ABV24631 (1-2452)
 OY 1 MetyaglnleuHlspProalaLeuGluThrAlaMetThrLeuPheProValLeu 20
 DB 197 ATGAACAATACTTCTTCCTGCGAACCACCTCAATGACATTTATCCAGTGTG 256
 OY 21 LeupheValaIaGlyLeuLeuProSerPheProAlaAsnGluAspLysAspProAla 40

DB 257 TTTCTCTGGTGTCTGGGCTGCTTCATCTTTTCCAGCAATGAAATAGATCCCGCT 316
 OY 41 PheThrAlaLeuLeuThrThrGlnThrGlnValaGlnArgGluLeuValaAsnLysHsAsn 60
 DB 317 TTACTGCTTTGTTAAACCAACCAACCAAGCAAGCAAGCAAGCAAGCAAGCAAGCAAG 376
 OY 61 GluLeuArgArgAlaValSerProProAlaArgAsnMetLeuLysMetGluTrpAsnLys 80
 DB 377 GAACGTAGAGAGACAGTATCTCCCTCCAGCAAAATGCTGAAAGATGAAAGCAAA 436
 OY 81 GluAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnTrpArgHsSerAsnPro 100
 DB 437 GAGCTGCAGCAAAATGCCCAAAAGTGGCAAAACCAATGCAATTCAGACAGTAACCA 496
 OY 101 LysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTyrMetSerAlaProSer 120
 DB 497 AAGGATCGAATGACAAAGTCTAAATGCTGAGAAATCTTACTGTCAAGTCTCCAGC 556
 OY 121 SerTrpSerGlnAlaAlaGlnSerTrpPheAspGluTyrAsnAspPheAspPheGlyVal 140
 DB 557 TCATGCTCACAAGCAATCCAAAGCTGTTGATGAGTACAAATGATTTGACTTGGTGA 616
 OY 141 GlyProLysThrProAsnAlaValaGlyHsTrpThrGlnValaValTrpTyrSerSer 160
 DB 617 GGCCCAAGACTCCCAACCCAGTGTGGACATTTATACACAGGTGTTGGTACTCTTCA 676
 OY 161 TyrLeuValaGlyCysGlyAsnAlaTyrCysProAsnGlnLysValaLeuLysTrpTyr 180
 DB 677 TACCTGCTTGATGTGGAAATCCCTACTCTCCCAATCAAAAGTCTTAAATACTACTAT 736
 OY 181 ValCysGlnTyrCysProAlaGlyAsnTrpAlaAsnArgLeuTyrValProTyrGluGln 200
 DB 737 GTTCCCAATATTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 796
 OY 201 GylAlaProCysAlaSerCysProAsnAsnCysAspAspGlyLeuLysThrAsnGlyCys 220
 DB 797 GGAGCACTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 856
 OY 221 LysTyrGluAspLeuTyrSerAsnCysLysSerLeuLysLeuThrCysLysHs 240
 DB 857 AAGTACGAAGACTCTATAGTACTGTAAGCTTGAAGCTTGAAGCTTGAAGCTTGAAGCTTGA 916
 OY 241 GlnLeuValaArgAspSerCysLysAlaSerCysAsnCysSerAsnSerIleTyr 258
 DB 917 CAGTTGTCAGAGGACAGTTCGCAAGGCTCTGCAATTTGTCAAAACAGCATTTAT 970
 RESULT 7
 ID ABV25272 standard; cDNA: 2452 BP.
 XX
 AC ABV25272;
 XX
 DT 16-SEP-2002 (first entry)
 XX
 DE Human prostate expression marker cDNA 25263.
 XX
 KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
 KW pharmacogenomic marker; gene; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200160860-A2.
 XX
 PD 23-AUG-2001.
 XX
 PF 20-FEB-2001; 2001WO-US05171.
 XX
 PR 17-FEB-2000; 2000US-183319P.
 PR 16-MAR-2000; 2000US-189862P.
 PR 25-MAY-2000; 2000US-207454P.
 PR 09-JUN-2000; 2000US-211314P.
 PR 18-JUL-2000; 2000US-219007P.

PR 13-DEC-2000; 2000US-255281P.
XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
PI Schlegel R, Endege WO, Monahan JE;
XX MPI: 2001-662795/76.
XX
XX Novel isolated nucleic acid molecule associated with cancerous state of
PT prostate cells and correlating with presence of prostate cancer, useful
PT for detecting presence of prostate cancer, stage of prostate cancer
XX
PS Claim 1: Page 4961; 11750pp; English.

CC The invention relates to an isolated nucleic acid molecule (I) comprising
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
CC specification or its complement. (I) is useful for:
CC (a) assessing whether a patient is afflicted with prostate cancer;
CC (b) monitoring the progression of prostate cancer in a patient;
CC (c) assessing the efficacy of a test compound to inhibit prostate
CC cancer in a patient;
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer
CC in a patient;
CC (e) selecting a composition for inhibiting prostate cancer in a patient;
CC (f) assessing the prostate cell carcinogenic potential of a compound;
CC (g) determining whether prostate cancer has metastasized in a patient;
CC (h) assessing the aggressiveness or indolence of prostate cancer in a
CC patient;
CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.

SQ Sequence 2452 BP; 787 A; 467 C; 444 G; 754 T; 0 other;

Alignment Scores:
Pred. No.: 1,43e-135 Length: 2452
Score: 1428.00 Matches: 257
Percent Similarity: 99.61% Conservative: 0
Best Local Similarity: 99.61% Mismatches: 1
Query Match: 99.44% Indels: 0
Gaps: 0

US-09-698-781-3 (1-258) x ABV25272 (1-2452)

OY 1 MetysGlnIleuHisProAlaLeuGluThrAlaMetThrLeuPheProValIleu 20
DB 197 ATGAAACAAATCTTCTCTCTGCTGGAACCTGCAATGACATTTATCCCGTGTG 256
OY 21 LeuPheLeuValAlaGlyLeuLeuProSerPheProAlaAsnGluAspLysAspProAla 40
DB 257 TTGCTCTGGTGGTGGCTGCTCCATCTTTCCACCAATGAAATGAAGATCCCGCT 316
OY 41 PheThrAlaLeuLeuThrThrGlnThrGlnValGlnArgGluIleValAsnHisAsn 60
DB 317 TTTACGTCTTGTAACTACCCCAACCAAGTGAAGGAGATGTGAATTAACCAAT 376
OY 61 GluLeuArgAlaValSerProAlaArgAsnMetLeuLysMetGluThrAspLys 80
DB 377 GAATGAGAGAGAGATATCTCCCTGCCAAGAACATGCTGAAGATGGAATGAAACAA 436
OY 81 GluAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnLysArgHisSerAsnPro 100
DB 437 GAGGCTCAGCAAAATGCCCAAAAGTGGCAAAACAGTGCATATACAGACACAGTAACCA 496
OY 101 LysAspArgMetThrSerLeuLysCysGlyGluAsnLeuLysMetSerSerAlaProSer 120
DB 497 AAGGATGGAATGACAAATCTAAATGTGTGAGATCTTCAATGATCAAGTCCCTCAGC 556
OY 121 SerTrpSerGlnAlaIleGlnSerTrpPheAspGluTrpAsnAspPheAspPheGlyVal 140
DB 557 TCAATGTCACAGCAATCCAAAGCTGGTTGATGATGATCAATGATTTTGACTTTGGGTGA 616
OY 141 GlyProLysThrProAsnAlaValAlaGlyHisLysThrGlnValAlaLysTrpSerSer 160
DB 617 GGGCAAAAGACTCCCAAGCAGGTGGTGGACATATATACACAGGTGTTGGTACTCTCA 676

OY 161 TyrLeuValGlyCysGlnAsnAlaTyrCysProAsnGlnLysValLeuLysTyrTyr 180
DB 677 TACCTGCTTGATGTGGAATGCTACTGTCCCAATCAAAAAGTTCAAAATACTACTAT 736
OY 181 ValCysGlnTrpCysProAlaGlnAsnTrpAlaAsnArgLeuTyrValProTyrGluGln 200
DB 737 GTTTCGCAATATATGTCTCTGCTGGAATGTGGCTAATGACTATATATGCCCTATGAACAA 796
OY 201 GlyAlaProCysAlaSerCysProAspAsnCysAspAspGlyLeuCysThrAsnGlyCys 220
DB 797 GGAGCACTTGTGCTCCAGTTCCCAAGATGATGATGATGATGATGATGATGATGATGATG 856
OY 221 LysTrpGlnAspLeuTyrSerAsnCysLysSerLeuLysLeuThrCysLysHis 240
DB 857 AAGTACGACAGATCTCTATGATGATGATGATGATGATGATGATGATGATGATGATGAT 916
OY 241 GlnLeuValArgAspSerCysLysAlaSerCysAsnCysSerAsnSerIleTyr 258
DB 917 CAGTGTGACGAGGACAGTGTGCAAGGCTCTGCAATGTTCAAACACATTTAT 970

RESULT 8
ABV25706
ID ABV25706 standard; cDNA; 2452 BP.
XX
XX ABV25706:
AC
XX
DT 16-SEP-2002 (first entry)
XX
XX Human prostate expression marker cDNA 25697.
DE
XX
XX Human prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
KM pharmacogenomic marker; gene; ss.
KW
XX Homo sapiens.
XX
XX MO200160860-A2.
XX
XX 23-AUG-2001.
PD
XX
XX 20-FEB-2001; 2001MO-US05171.
PE
XX
XX 17-FEB-2000; 2000US-183319P.
PR 16-MAR-2000; 2000US-189862P.
PR 25-MAY-2000; 2000US-207454P.
PR 09-JUN-2000; 2000US-211314P.
PR 18-JUL-2000; 2000US-219007P.
PR 13-DEC-2000; 2000US-255281P.
XX
XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
PA
XX Schlegel R, Endege WO, Monahan JE;
PI
XX MPI: 2001-662795/76.
DR
XX Novel isolated nucleic acid molecule associated with cancerous state of
PT prostate cells and correlating with presence of prostate cancer, useful
PT for detecting presence of prostate cancer, stage of prostate cancer
XX
PS Claim 1: Page 5156-5157; 11750pp; English.

CC The invention relates to an isolated nucleic acid molecule (I) comprising
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
CC specification or its complement. (I) is useful for:
CC (a) assessing whether a patient is afflicted with prostate cancer;
CC (b) monitoring the progression of prostate cancer in a patient;
CC (c) assessing the efficacy of a test compound to inhibit prostate
CC cancer in a patient;
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer
CC in a patient;
CC (e) selecting a composition for inhibiting prostate cancer in a patient;
CC (f) assessing the prostate cell carcinogenic potential of a compound;
CC (g) determining whether prostate cancer has metastasized in a patient;

CC (h) assessing the aggressiveness or indolence of prostate cancer in a
 CC patient;
 CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.
 XX
 SO Sequence 2452 BP; 787 A; 467 C; 444 G; 754 T; 0 other;

Alignment Scores:

Pred. No.:	1,43e-135	Length:	2452
Score:	1428.00	Matches:	257
Percent Similarity:	99.61%	Conservative:	0
Best Local Similarity:	99.61%	Mismatches:	1
Query Match:	99.44%	Indels:	0
DB:	23	Gaps:	0

US-09-698-781-3 (1-258) x ABV25706 (1-2452)

```

OY 1 MettysGlnlleuHisProalaLeuGluThrThraIaMetThleupheProvalleu 20
DB 197 ATGAACAATAATCTTCATCTCTGGAACCACTGCATGACATATATCCAGTGTG 256
OY 21 LeupheleuValaIaGlyLeuLeuProserPheProalaasnGluasplysaproala 40
DB 257 TTGTCCTGCTGCTGGGCTGCTTCATCTTTCCAGCAATGAAGATAGATCCGCT 316
OY 41 PheThraIaLeuLeuThrThrGlnThrGlnValaGlnArgGluIleValaAsnLysHisasn 60
DB 317 TTACTGCTTTGTAAACCAACCAACAAACAAAGGAGGAGATGTGAATAGACAAAT 376
OY 61 GlutauArgArgAlaValSerProProalaargAsnMetLeuLysMetCylurpsnLys 80
DB 377 GAACGAGAGAGAGCAATCTCCCTGCGAACAACATGCTAAAGTGAAGGAAACAA 436
OY 81 GluaIaAlaIaIaasnAlaGlnLysTrpAlaasnGlnCysAsnTyArgHisserasnPro 100
DB 437 GAGCGTGACGAATGCCCAAAAGTGGGCAAAACAGTGCAATACAGACAGTAAACCA 496
OY 101 LysAspArgMetThrSerLeuLysCysGlyGluasnLeuTyMetSerSerAlaProser 120
DB 497 AAGGATCAATGACAAAGCTTAATAATGTGTGAATCTCTACATGTCAAGTCCCTCACG 556
OY 121 SerTPSerGlnAlaIleGlnSerTrpPheaspGluTyArgAsnAspPheaspheGlyVal 140
DB 557 TCATGCTCAACAAGCAATCCAAAGCTGTTGATGATACATGATTTGATTTGGTGA 616
OY 141 GlyProLysThrProasnAlaValaIaGlyHisTyThrGlnValaIaTrpTyrSerSer 160
DB 617 GGGCCAAAGACTCCCAACGACAGTGTGACATATATACACAGTGTGTTGTAAGTCTTCA 676
OY 161 TytleuValaIaGlyCysGlyAsnAlaTyrcysProasnGlnLysValaLeuLysTyrrtyr 180
DB 677 TACCTCGTTGATGTGGAATGCCACTGTCCCAATCAAAAAGTCTTAATAACACTAT 736
OY 181 ValCysGlnTyrcysProalaGlyasnTrpAlaasnArgLeuTyValProtyrGlnGln 200
DB 737 GTTTCACAAATATGTCTGCTGCTGTAATGGCTAATAGACTATATGCTTATGACAA 796
OY 201 GlyAlaProCysAlaSerCysProaspasnCysAspAspGlyLeuLysThrAsnGlyCys 220
DB 797 GGAGACCTTGTGCGAGTGTGCCAATATGCTGACGATGAGTATGACCAATGATGTTTC 856
OY 221 LysTyrgLysAspLeuTySerasnCysLysSerLeuLysLeuThrLeuThrcysLysHis 240
DB 857 AAGTACGAAGATCTTATATGTAAGTAAAGTTTGAAGCTCACAATTAACCTGTAACAT 916
OY 241 GlnleuValaIaArgAspSerCysLysAlaSerCysAsnCysSerAsnSerIleTyrr 258
DB 917 CAGTGTGTCAGGAGACATTGCAAGGCTCTCTCAATTTGTTCAACAGCATTTAT 970

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RESULT 9
 ABV28467
 ID ABV28467 standard; cDNA: 2452 BP.
 XX
 AC ABV28467;

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XX 16-SEP-2002 (first entry)
DT Human prostate expression marker cDNA 28458.
XX
DE Human: prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
XX pharmacogenomic marker; gene; ss.
XX Homo sapiens.
OS WO200160860-A2.
XX
PN 23-AUG-2001.
XX
PD 20-FEB-2001; 2001WO-US05171.
XX
PF 17-FEB-2000; 2000US-183319P.
XX
PR 16-MAR-2000; 2000US-189862P.
XX
PR 25-MAY-2000; 2000US-207454P.
XX
PR 09-JUN-2000; 2000US-211314P.
XX
PR 18-JUL-2000; 2000US-219007P.
XX
PR 13-DEC-2000; 2000US-255281P.
XX
PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
XX
PI Schlegel R, Endege WO, Monahan JE;
XX
PI WPI: 2001-662795/76.
XX
DR Novel isolated nucleic acid molecule associated with cancerous state of
XX prostate cells and correlating with presence of prostate cancer, useful
XX for detecting presence of prostate cancer, stage of prostate cancer -
XX
XX Claim 1: Page 5942-5943; 11750pp; English.
XX
XX The invention relates to an isolated nucleic acid molecule (I) comprising
XX a nucleotide sequence given in Tables 1-9 (ABV0010-ABV62213) of the
XX specification or its complement. (I) is useful for:
XX (a) assessing whether a patient is afflicted with prostate cancer;
XX (b) monitoring the progression of prostate cancer in a patient;
XX (c) assessing the efficacy of a test compound to inhibit prostate
XX cancer in a patient;
XX (d) assessing the efficacy of a therapy for inhibiting prostate cancer
XX in a patient;
XX (e) selecting a composition for inhibiting prostate cancer in a patient;
XX (f) assessing the prostate cell carcinogenic potential of a compound;
XX (g) determining whether prostate cancer has metastasized in a patient;
XX (h) assessing the aggressiveness or indolence of prostate cancer in a
XX patient;
XX (i) is also useful as a pharmacodynamic or pharmacogenomic marker.
XX
XX Sequence 2452 BP; 787 A; 467 C; 444 G; 754 T; 0 other;
XX
XX Alignment Scores:
XX Pred. No.: 1,43e-135 Length: 2452
XX Score: 1428.00 Matches: 257
XX Percent Similarity: 99.61% Conservative: 0
XX Best Local Similarity: 99.61% Mismatches: 1
XX Query Match: 99.44% Indels: 0
XX DB: 23 Gaps: 0
XX
XX US-09-698-781-3 (1-258) x ABV28467 (1-2452)
XX
XX 1 MettysGlnlleuHisProalaLeuGluThrThraIaMetThleupheProvalleu 20
DB 197 ATGAACAATAATCTTCATCTCTGGAACCACTGCATGACATATATCCAGTGTG 256
OY 21 LeupheleuValaIaGlyLeuLeuProserPheProalaasnGluasplysaproala 40
DB 257 TTGTCCTGCTGCTGGGCTGCTTCATCTTTCCAGCAATGAAGATAGATCCGCT 316
OY 41 PheThraIaLeuLeuThrThrGlnThrGlnValaGlnArgGluIleValaAsnLysHisasn 60
DB 317 TTACTGCTTTGTAAACCAACCAACAAACAAAGGAGGAGATGTGAATAGACAAAT 376

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Db 317 TTTACTGCTTTGTTAAACCAACCAAGTCGAAAGGAGATGTGTAATGAACACAAAT 376
 Qy 61 GtluLeuAArgAlaValSerProAlaArgAsnMetLeuYsMetGtUTPAsnLys 80
 Db 377 GAACTGAGGACGACGATCTCCCTGCGAGAACATGCTGAAGATGGAATGGAAACAAA 436
 Qy 81 GtAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnTyrArgHisSerAsnPro 100
 Db 437 GAGGCTGACCAAAATGCCCAAAAGTGGGCAACACAGTCGATTAACAGACAGTAACCCA 496
 Qy 101 LysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTyrMetSerSerAlaProSer 120
 Db 497 AAGCATCGAATGACAAAGTCTAAATGCGTGAGAAATCTCTACATGTCGAACTGCCTCCAGC 556
 Qy 121 SerTrpSerGlnAlaIleGlnSerTrpPheAspGluTyrAsnAspPheAspPheGlyVal 140
 Db 557 TCAATGGTCACAGCAACATCCAAAGCTGGTTGATGAGTACAAATGATTTTGACTTGGGTGA 616
 Qy 141 GlyProLysThrProAsnAlaValAlaGlyHisTyrThrGlnValAlaTyrPtyrSerSer 160
 Db 617 GGGCAAAAGACTCCCAACGCAAGTGGTGGACATATATACACAGGTTGTGGTACTCTCA 676
 Qy 161 TyrLeuValGlyCysGlyAsnAlaTyrCysProAsnGlnLysValLeuLysTyrTyr 180
 Db 677 TACCTCGTGGATGTGAAATGCTCTACTGTCCCAATCAAAAGTCTTAAATACTACTAT 736
 Qy 181 ValCysGlnTyrCysProAlaGlyAsnTrpAlaAsnArgLeuTyrValProTyrGluGln 200
 Db 737 GTTTCACCAATATTTGCTCTGCTGTAATGGCTATATAGACTATATATGCTTATGAAACAA 796
 Qy 201 GlyAlaProCysAlaSerCysProAspAsnCysAspAspGlyLeuCysTrpHisGlyCys 220
 Db 797 GAGACACTTGTGGCACTTGGCCAGATTAAGTGCAGATGAGCATGAGCAATGAGTGGTGC 856
 Qy 221 LysTyrGlnAspLeuTyrSerAsnCysLysSerLeuLysLeuThrLeuThrCysLysHis 240
 Db 857 AAGTACGAAATCTCTATTAATTAATGTTGAAGCTCACATTAACCTGTAAACAT 916
 Qy 241 GtluLeuValArgAspSerCysLysAlaSerCysAsnCysSerAsnSerLysTyr 258
 Db 917 CAGTGGTCAAGGACAGTCTCAAGGCTCTCCGCAATGTTCAAAACACATTTAT 970
 RESULT 10
 ABV28648 standard: cDNA: 2452 BP.
 ID ABV28648: standard: cDNA: 2452 BP.
 AC ABV28648:
 XX 16-SEP-2002 (first entry)
 DE Human prostate expression marker cDNA 28639.
 KW Human: prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
 KW pharmacogenomic marker; gene; ss.
 XX Homo sapiens.
 XX W0200160860-A2.
 FN 23-AUG-2001.
 PD 20-FEB-2001: 2001WO-US05171.
 XX 17-FEB-2000: 2000US-183319P.
 PR 16-MAR-2000: 2000US-189862P.
 PR 25-MAY-2000: 2000US-207454P.
 PR 09-JUN-2000: 2000US-211314P.
 PR 18-JUL-2000: 2000US-219007P.
 PR 13-DEC-2000: 2000US-235281P.
 XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
 PA Schlegel R, Endege WO, Monahan JE.
 XX PI

XX WPI: 2001-662795/76.
 DR Novel isolated nucleic acid molecule associated with cancerous state of
 XX prostate cells and correlating with presence of prostate cancer, useful
 PT for detecting presence of prostate cancer, stage of prostate cancer
 PT
 PS Claim 1: Page 6006; 11750pp; English.
 XX
 CC The invention relates to an isolated nucleic acid molecule (1) comprising
 CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
 CC specification or its complement. (1) is useful for:
 CC (a) assessing whether a patient is afflicted with prostate cancer;
 CC (b) monitoring the progression of prostate cancer in a patient;
 CC (c) assessing the efficacy of a test compound to inhibit prostate
 CC cancer in a patient;
 CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer
 CC in a patient;
 CC (e) selecting a composition for inhibiting prostate cancer in a patient;
 CC (f) assessing the prostate cell carcinogenic potential of a compound;
 CC (g) determining whether prostate cancer has metastasized in a patient;
 CC (h) assessing the aggressiveness or indolence of prostate cancer in a
 CC patient;
 CC (1) is also useful as a pharmacodynamic or pharmacogenomic marker.
 XX
 SQ Sequence 2452 BP; 787 A; 467 C; 444 G; 754 T; 0 other;
 Alignment Scores:
 Pred. No.: 1,43e-135 Length: 2452
 Score: 1428.00 Matches: 257
 Percent Similarity: 99.61% Conservative: 0
 Best Local Similarity: 99.61% Mismatches: 1
 Query Match: 99.44% Indels: 0
 DB: 23 Gaps: 0
 US-09-698-781-3 (1-258) x ABV28648 (1-2452)
 Qy 1 MetLysGlnIleLeuHisProAlaLeuGluThrAlaMetThrLeuPheProValLeu 20
 Db 197 ATGAACCAATACTTCACTCCGCTCTGGAACCACTGCAATGACATTAATCCAGTCTG 256
 Qy 21 LeuPheLeuValAlaGlyLeuLeuProSerPheProAlaAsnGluAspLysAspProAla 40
 Db 257 TTGTTCTGCTGGTGGCTGCTCTTCCATCTTTCACAGAAATGAAGATTAAGATCCGCT 316
 Qy 41 PheThrAlaLeuLeuThrThrGlnThrGlnValGlnArgGluIleValAsnLysHisAsn 60
 Db 317 TTTACTGCTTTGTTAACCACCCCAACACAACTGCAGAAAGGAGATGTGAATTAACCCACAT 376
 Qy 61 GtluLeuArgArgAlaValSerProAlaArgAsnMetLeuYsMetGtUTPAsnLys 80
 Db 377 GAACTGAGGACGACGATCTCCCTGCGAGAACATGCTGAAGATGGAATGGAAACAAA 436
 Qy 81 GtAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnTyrArgHisSerAsnPro 100
 Db 437 GAGGCTGACCAAAATGCCCAAAAGTGGGCAACACAGTCGATTAACAGACAGTAACCCA 496
 Qy 101 LysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTyrMetSerSerAlaProSer 120
 Db 497 AAGCATCGAATGACAAAGTCTAAATGCGTGAGAAATCTCTACATGTCGAACTGCCTCCAGC 556
 Qy 121 SerTrpSerGlnAlaIleGlnSerTrpPheAspGluTyrAsnAspPheAspPheGlyVal 140
 Db 557 TCAATGGTCACAGCAACATCCAAAGCTGGTTGATGAGTACAAATGATTTTGACTTGGGTGA 616
 Qy 141 GlyProLysThrProAsnAlaValAlaGlyHisTyrThrGlnValAlaTyrPtyrSerSer 160
 Db 617 GGGCAAAAGACTCCCAACGCAAGTGGTGGACATATATACACAGGTTGTGGTACTCTCA 676
 Qy 161 TyrLeuValGlyCysGlyAsnAlaTyrCysProAsnGlnLysValLeuLysTyrTyr 180
 Db 677 TACCTCGTGGATGTGAAATGCTCTACTGTCCCAATCAAAAGTCTTAAATACTACTAT 736

OY 181 ValysgIntYrCysPrioAlaGlyAsnTrpAlaAsnArgLeuTyrValProTyrGluGln 200
 DB 737 GTTGGCCAAATATTGCTCCGTGTGTAATGGCTAATAGACTAATGTCCTATATGACAA 796
 OY 201 GllValAProCysAlaSerCysPrioAspAsnCysAspAspGlyLeuCysThrAsnGlyys 220
 DB 797 GGAGCACCCTTGTCCAGTGGCCAGATGATGACGATGACATGACCAATGGTTGC 856
 OY 221 LysTyrGluAspLeuTyrSerAsnCysLysSerLeuLysLeuThrLeuThrCysLysHis 240
 DB 857 AAGTACGAAAGATCTCTAATAGTAAAGTTTGAAGCTCACCATTAACTGTAACAT 916
 OY 241 GlnLeuValArgAspSerCysLysAlaSerCysAsnCysSerAsnSerIleTyr 258
 DB 917 CAGTTGGTCAGGACGACAGTTCAGAGCCCTCTGCAATTGTTCAACACAGATTAT 970
 RESULT 11
 ABL67806
 ID ABL67806 standard; DNA; 2128 bp.
 XX
 AC ABL67806;
 XX
 DT 15-MAY-2002 (first entry)
 XX
 DE Oesophagus cancer related gene sequence SEQ ID NO: 6143.
 XX
 KM Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
 KM stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
 KM cytosaratic; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma;
 KM gene; ds.
 XX
 OS Homo sapiens.
 XX
 PN W0200194629-A2.
 PD 13-DEC-2001.
 PF 30-MAY-2001; 2001MO-US10838.
 XX
 PR 05-JUN-2000; 2000US-209473P.
 PR 05-JUN-2000; 2000US-209531P.
 PR 18-SEP-2000; 2000US-231133P.
 PR 18-SEP-2000; 2000US-233617P.
 PR 20-SEP-2000; 2000US-234009P.
 PR 20-SEP-2000; 2000US-234034P.
 PR 20-SEP-2000; 2000US-234052P.
 PR 22-SEP-2000; 2000US-234509P.
 PR 22-SEP-2000; 2000US-234567P.
 PR 25-SEP-2000; 2000US-234923P.
 PR 25-SEP-2000; 2000US-234924P.
 PR 25-SEP-2000; 2000US-235077P.
 PR 25-SEP-2000; 2000US-235082P.
 PR 25-SEP-2000; 2000US-235134P.
 PR 25-SEP-2000; 2000US-235280P.
 PR 26-SEP-2000; 2000US-235637P.
 PR 26-SEP-2000; 2000US-235638P.
 PR 27-SEP-2000; 2000US-235711P.
 PR 27-SEP-2000; 2000US-235720P.
 PR 27-SEP-2000; 2000US-235840P.
 PR 27-SEP-2000; 2000US-235863P.
 PR 28-SEP-2000; 2000US-236028P.
 PR 28-SEP-2000; 2000US-236032P.
 PR 28-SEP-2000; 2000US-236033P.
 PR 28-SEP-2000; 2000US-236034P.
 PR 28-SEP-2000; 2000US-236033P.
 PR 28-SEP-2000; 2000US-236109P.
 PR 28-SEP-2000; 2000US-236111P.
 PR 29-SEP-2000; 2000US-236842P.
 PR 29-SEP-2000; 2000US-236891P.
 PR 02-OCT-2000; 2000US-237172P.
 PR 02-OCT-2000; 2000US-237173P.
 PR 02-OCT-2000; 2000US-237278P.
 PR 02-OCT-2000; 2000US-237294P.
 PR 02-OCT-2000; 2000US-237295P.
 PR 02-OCT-2000; 2000US-237295P.

PR 02-OCT-2000; 2000US-237316P.
 PR 03-OCT-2000; 2000US-237425P.
 PR 03-OCT-2000; 2000US-237598P.
 PR 03-OCT-2000; 2000US-237604P.
 PR 03-OCT-2000; 2000US-237606P.
 PR 03-OCT-2000; 2000US-237608P.
 PR 01-NOV-2000; 2000US-244867P.
 PR 01-NOV-2000; 2000US-245084P.
 XX
 PA (AVAL-) AVALON PHARM.
 XX
 PI Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;
 PI Soppet DR, Weaver Z;
 XX
 DR WPI: 2002-188264/24.
 XX
 PT Screening for anti-neoplastic agent involves exposing cells to a
 PT chemical agent to be tested for anti-neoplastic activity, and
 PT determining a change in expression of a gene of a signature gene set -
 PS
 PS Claim 1; SEQ ID 6143; 44pp; English.
 XX
 CC The present invention describes a method (M1) for screening for an
 CC anti-neoplastic agent. The method involves exposing cells to a chemical
 CC agent to be tested for anti-neoplastic activity, determining a change in
 CC expression of at least one gene (I) of a signature gene set, where (I)
 CC comprises a sequence (S) selected from 8447 sequences (given in ABL61664
 CC to ABL70110), or is at least 95% identical to (S), where a change in
 CC expression is indicative of anti-neoplastic activity. (I) has cytostatic
 CC activity and can be used in gene therapy. M1 can be used for screening
 CC an anti-neoplastic agent, and can be used for producing a product which
 CC is the data collected with respect to the anti-neoplastic agent as a
 CC result of M1, and the data is sufficient to convey the chemical
 CC structure and/or properties of the agent. M1 can be used in the
 CC treatment of cancer such as colon, breast, stomach, lung, thyroid,
 CC oesophageal, ovarian, kidney, prostate or pancreatic cancer,
 CC adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer,
 CC infiltrating lobular cancer, squamous cell carcinoma, neuroendocrine
 CC carcinoma, papillary carcinoma and Wilm's tumour.
 XX
 SQ Sequence 2128 bp; 734 A; 397 C; 380 G; 617 T; 0 other;
 Alignment Scores:
 Pred. No.: 2,26e-131 Length: 2128
 Score: 1386.00 Matches: 249
 Percent Similarity: 99.60% Conservative: 0
 Best Local Similarity: 99.60% Mismatches: 1
 Query Match: 96.52% Indels: 0
 DB: 24 Gaps: 0
 US-09-698-781-3 (1-258) x ABL67806 (1-2128)
 OY 9 LeuGluTrpThrAlaMetThrLeuPheProValLeuLeuValAlaGlyLeuLeu 28
 DB 1 CTGGAAACCATGCAATGACATTAATTCACAGTGGTGTGCTGGGTGGGCTCTT 60
 OY 29 ProSerPheProAlaAsnGluAspLysAspProAlaPheThrAlaLeuLeuThrGln 48
 DB 61 CCATCTTTCCAGCAAAATGAAGATAGGATCCCTTTTACTGTTGTTTACACCCAA 120
 OY 49 ThrGlnValGlnArgGluIleValAlaAsnLysHisAsnGluLeuArgAlaValSerPro 68
 DB 121 ACACAGTGCAGAAAGGAGATTGTAATAGCAATGACAGAGAGAGAGAGATCTCC 180
 OY 69 ProAlaArgAsnMetLeuLysMetGluTrpAsnLysGluAlaAlaAsnAlaGlnLys 88
 DB 181 CCTCCGGAACATGCTGAGATGAAATGGAACAAAGAGGTGAGCAAAAGCCCAAAAG 240
 OY 89 TrpAlaAsnGlnCysAsnTyrArgHisSerAsnProLysAspArgMetThrSerLeuLys 108
 DB 241 TGGGCAAAACGATGCAATTACAGACAGTAAACCAAGATGCAATGACAGATCTAAA 300
 OY 109 CysGlyGluAsnLeuTyrMetSerAlaProSerSerTrpSerGlnAlaIleGlnSer 128

Db 301 TGTGGGAGAAATCTACTACTGTAAGTGGCTCCAGCTACTGTCACAGCAATCCAAAGC 360
 Oy 129 TTPPheaspGluTyrAsnAspPheaspPheGlyValGlyProLysThrProAsnAlaVal 148
 Db 361 TGGTTGATGAGTACAAATGATTTGACTTGTGTAGGGCCAAAGACTCCCAACGAGTG 420
 Oy 149 ValGlyHisThrGlnValValTrrPyrSerSerTyrLeuValGlyCysGlyAsnAla 168
 Db 421 GTTGGACATTATACACAGGTGTTGGTACTCTTCATACCTCGTGGATGGAAATGCC 480
 Oy 169 TTYCYPProAsnGlnLysValLeuLysTyrTyrTyrValCysGlnTyrCysProAlaGly 188
 Db 481 TACTGTCCCAATCAAAAGTCTTAAATCTACTATGTTGGCAATATTTGCTGCGGT 540
 Oy 189 AsnTrrPalaAsnArgLeuTyrValProTyrGluGlnGlyAlaProCysAlaSerCysPro 208
 Db 541 AATTGGGCTAATAGACTATATGTCCCTTATGACAAGAGACACCTTGTGCCAGTTGCCCA 600
 Oy 209 AspAsnCysAspAspGlyLeuCysThrAsnGlyCysLysTyrGluAspLeuTyrSerAsn 228
 Db 601 GATTAAGTGTGACATGACATGACATGACCAATAGTTGCAAGTACGAAATCTCTATAGTAC 660
 Oy 229 CysLysSerLeuLysLeuThrLeuThrCysLysHisGlnLeuValArgAspSerCysLys 248
 Db 661 TGTAAAGTTTGAAGCTCAGCATTAACCTGTAAACATCAGTTGGTCAGGAGAGTTGCAAG 720
 Oy 249 AlaSerCysAsnCysSerAsnSerIleTyr 258
 Db 721 GCCTCGTCGAATTTGTTCAACACAGCATTTAT 750
 RESULT 12
 AAS70843 standard: cDNA: 1386 BP.
 AAS70843:
 13-FEB-2002 (first entry)
 DNA encoding novel human diagnostic protein #6647.
 Human: chromosome mapping: gene mapping: gene therapy: forensic;
 food supplement: medical imaging; diagnostic; genetic disorder: ss.
 Homo sapiens.
 MO200175067-A2.
 11-OCT-2001.
 30-MAR-2001: 2001MO-US08631.
 31-MAR-2000: 2000US-0540217.
 23-AUG-2000: 2000US-0649167.
 (HYSE-) HYSEQ INC.
 Drmanac RT, Liu C, Tang YT;
 WPI: 2001-639362/73.
 P-PSDB: ABG06656.
 New isolated polynucleotide and encoded polypeptides: useful in
 diagnostics, forensics, gene mapping, identification of mutations
 responsible for genetic disorders or other traits and to assess
 biodiversity
 Claim 1: SEQ ID NO 6647; 103pp; English.
 The invention relates to isolated polynucleotide (I) and
 polypeptide (II) sequences. (I) is useful as hybridisation probes,
 polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 and gene mapping, and in recombinant production of (II). The

CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at http://wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 1386 BP; 438 A; 293 C; 289 G; 366 T; 0 other;
 Alignment Scores:
 Pred. No.: 1,94e-92 Length: 1386
 Score: 1002.00 Matches: 184
 Percent Similarity: 80.84% Conservative: 27
 Best Local Similarity: 70.50% Mismatches: 45
 Query Match: 69.78% Indels: 5
 DB: 23 Gaps: 3
 US-09-698-781-3 (1-258) x AAS70843 (1-1386)
 Oy 1 MetLysGlnLeuHisProAlaLeuGluThr-Thr-----AlaMetThrLeuPhePyr 18
 Db 195 ATTAAGTACATATTTCTCTCTCTCAGAAAACCAACATTTCCAGCAATGGCTTACTACACC 254
 Oy 18 OValLeuLeuPheLeuValAlaGlyLeuLeuProSerPheProAlaAsnGluAspLysAs 38
 Db 255 GGTG---TTGTTCTGTGTTACTGTGCTGCTCCATCTTACTCTCA---GAAGCAAGAAGA 308
 Oy 38 ProAlaPheThrAlaLeuLeuThrThrGlnThrGlnValGlnArgGluLeuValAsnLys 58
 Db 309 TCCCGCTTTTACTCTCTTGTGTTAACCCAGCTTCAAGTCAAGAGGAGATGTAATAA 368
 Oy 58 SHAsnGluLeuArgArgAlaValAsnProProAlaArgAsnMetLeuLysMetGluTr 78
 Db 369 ACACAAATGAACCTTAAGGAAGACAGTCTCTCCAGCTCCAGTAACATGCTAAAGATGGAATG 428
 Oy 78 PAsnLysGluAlaAlaAlaAsnAlaGlnLysTrrPalaAsnGlnCysAsnTyrArgHisSe 98
 Db 429 GAGCAGAGAGGTATACCAAGATGCCCAAGAGTGGCAAAACAGTCACTTACCAACATAG 488
 Oy 98 RasnProLysAspArgMetThrSerLeuLysCysGlyGlnAsnLeuTyrMetSerSerAl 118
 Db 489 TGAATCCAGAGACCGCAAAACAGTACAGATGAGTGAATCTTATATGCAAGTGA 548
 Oy 118 AProSerSerTrrPserGlnAlaIleGlnSerTrrPheaspGluTyrAsnAspPheaspPh 138
 Db 549 CCTTACTTCTGCTCTTCTGTGAATCCAAAGCTGGTATACAGATCCAGATTTTGTGCTA 608
 Oy 138 eGlyValGlyProLysThrProAsnAlaValAlaGlyHisTyrThrGlnValValTrrPty 158
 Db 609 TGTGTAGAGACCAAAAGATGCCCAATGCGTGTGGACATATATCTAGCTTGTGGTA 668
 Oy 158 rSerSerTyrLeuValGlyCysGlyAsnAlaTyrCysProAsnGlnLysValLeuLysTyr 178
 Db 669 CTCACCTTACCAAGGTAGGCTGTGAATTTGCTACTGTCTCCATCAAGTACTCTAAATA 728
 Oy 178 rTyrTyrValCysGlnTyrCysProAlaGlyAsnTrrPalaAsnArgLeuTyrValProTy 198
 Db 729 CTACTATGTTGGCAATATTTCTCTGCTGTAATATATGATGAAGAATATACCCGCTA 788
 Oy 198 rGluGlnGlyAlaProCysAlaSerCysProAspAsnGlyLeuCysThrAs 218
 Db 789 CCACAAAGAACACCTTGTGCGGTGCTGATGACTGTACAAAGACATATGACACCA 848

XX 16-SEP-2002 (first entry)
 XX Human prostate expression marker CDNA 43205.
 DE Human prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
 KM pharmacogenomic marker; gene; ss.
 XX Homo sapiens.
 OS
 XX WO200160860-A2.
 XX
 XX PD 23-AUG-2001.
 XX
 XX PF 20-FEB-2001; 2001WO-US05171.
 XX
 XX PR 17-FEB-2000; 2000US-183319P.
 XX PR 16-MAR-2000; 2000US-189862P.
 XX PR 25-MAY-2000; 2000US-207454P.
 XX PR 09-JUN-2000; 2000US-211314P.
 XX PR 18-JUL-2000; 2000US-219007P.
 XX PR 13-DEC-2000; 2000US-255281P.
 XX
 XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
 XX
 XX PI Schlegel R, Endege WO, Monahan JE;
 XX
 XX DR WPI; 2001-662795/76.
 XX
 XX PT Novel isolated nucleic acid molecule associated with cancerous state of
 PT prostate cells and correlating with presence of prostate cancer, useful
 PT for detecting presence of prostate cancer, stage of prostate cancer -
 XX
 XX PS Claim 1; Page 8623; 11750pp; English.
 XX
 CC The invention relates to an isolated nucleic acid molecule (I) comprising
 CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
 CC specification or its complement. (I) is useful for:
 CC (a) assessing whether a patient is afflicted with prostate cancer;
 CC (b) monitoring the progression of prostate cancer in a patient;
 CC (c) assessing the efficacy of a test compound to inhibit prostate
 CC cancer in a patient;
 CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer
 CC in a patient;
 CC (e) selecting a composition for inhibiting prostate cancer in a patient;
 CC (f) assessing the prostate cell carcinogenic potential of a compound;
 CC (g) determining whether prostate cancer has metastasized in a patient;
 CC (h) assessing the aggressiveness or indolence of prostate cancer in a
 CC patient;
 CC (I) is also useful as a pharmacodynamic or pharmacogenomic marker.
 XX
 XX SQ Sequence 534 BP; 120 A; 127 C; 133 G; 153 T; 1 other:
 XX
 XX Alignment Scores:
 XX Pred. No.: 1, 87e-60 Length: 534
 XX Score: 683.00 Matches: 132
 XX Percent Similarity: 98.51% Conservative: 0
 XX Best Local Similarity: 98.51% Mismatches: 1
 XX Query Match: 47, 56% Indels: 1
 XX DB: 23 Gaps: 0
 XX
 XX US-09-698-781-3 (1-258) x ABV43214 (1-534)
 XX
 OY 1 MettysGinleleuHISProaLaLeuGluThrThraLaMetThrLeuPheProValLeu 20
 DB 434 ATGAACAATACTTCACTCTGCTGGAACCACTGCATGACATATTATCCAGTCTG 375
 OY 21 LeuPheLeuValAlaGlyLeuLeuProSerPheProLaLsnGluAspLysAspProLa 40
 DB 374 TTGTTCTGTGCTGGGCTGCTTCCATCTTTTCCAGCAATGAAGAATGATCCGCT 315
 OY 41 PheThraLaLeuLeuThrThGlnThGlnValGlnArgGluIleValAsnLysHisAsn 60
 |||||||

DB 314 TTACTGCTTTGTTAACCAACCAACCAAGTGCAGAAAGGAGATTGTGAATAAGCAAT 255
 OY 61 GluLeuArgArgAlaValSerProPoaLaArgAsnMetLeuLysMetGluTrpAsnLys 80
 DB 254 GAACGTAGAGAGACACTATCTCCCTGCCAGAACATCTGTAAGATGGAATGGAACAA 195
 OY 81 GluAlaAlaAlaAsnAlaGlnLysTrpAlaAsn-GlnCysAsnTrpArgHisSerAsnPr 100
 DB 194 GAGGCTGCAGCAATCCCAAAAGTGGCAACNCAGTGCATTAAGACAGACAGTAACCC 135
 OY 100 oLysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTrpMetSerSerAlaProse 120
 DB 134 AAGGATCGAATGCAGTCTAAATGTGTGAGAAATCTCATCATGTCAAGTCCCTCCAG 75
 OY 120 rSerTrpSerGlnAlaIleGlnSerTrpPheAspGluTrp 133
 DB 74 CTCATGTGCACAGCAATCCAAAGCTGTGTTGATGAGTAC 35
 RESULT 15
 AAS70842
 ID AAS70842 standard; CDNA: 683 BP.
 XX
 XX AAS70842;
 XX
 XX 13-FEB-2002 (first entry)
 XX
 XX DE DNA encoding novel human diagnostic protein #6646.
 XX
 XX KM Human; chromosome mapping; gene mapping; gene therapy; forensic;
 XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX OS Homo sapiens.
 XX
 XX PN WO200175067-A2.
 XX
 XX PD 11-OCT-2001.
 XX
 XX PF 30-MAR-2001; 2001WO-US08631.
 XX
 XX PR 31-MAR-2000; 2000US-0540217.
 XX PR 23-AUG-2000; 2000US-0649167.
 XX
 XX PA (HYSE-) HYSEQ INC.
 XX
 XX PI Drmanac RT, Liu C, Tang YF;
 XX
 XX DR WPI; 2001-639362/73.
 XX P-PSDB; ABG06655.
 XX
 XX PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX
 XX Claim 1; SEQ ID No 6646; 103pp; English.
 XX
 PS The invention relates to isolated polynucleotide (I) and
 XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human

CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pat_sequences.
XX

Sequence 683 BP; 215 A; 165 C; 141 G; 162 T; 0 other;

Alignment Scores:

Prod. No.:	2.98e-45	Length:	683
Score:	535.50	Matches:	119
Percent Similarity:	64.73%	Conservative:	15
Best Local Similarity:	57.49%	Mismatches:	31
Query Match:	37.29%	Indels:	44
DB:	23	Gaps:	4

US-09-698-781-3 (1-258) x AAST0842 (1-683)

```
OY 1 MetLysGlnLeuHisProAlaLeuGluThr-Thr-----AlaMetThrLeuPhePr 18
DB 177 ATAAAGTAGATATTTCATCTCTGCTCAGAAACACATTTCAGCAATGGCTTACTACC 236
OY 18 oValLeuLeuPheLeuValAlaGlyLeuLeuProSerPheProAlaAsnGlnAspLysAs 38
DB 237 GGTG---TTGTTCTGTGTTACTGCTGCTGCTCCATCTTACTGCA---GAAGGAAAGGA 290
OY 38 ProAlaPheThrAlaLeuLeuThrThrGlnThrGlnValGlnArgGluLeuValAsnLy 58
DB 291 TCCCGCTTTACTGTTTGTGTTACCCAGCCAGTTGCAAGTACAAAGGAGATGTAATAA 350
OY 58 sHisAsnGluLeuArgArgAlaValSerProAlaArgAsnMetLeuLysMetGluTr 78
DB 351 ACACAAATGAACTAGAGAAAGCACTCTCCACTGCCAGTACATGCTAAAGATGGAATG 410
OY 78 pAsnLysGlnAlaAlaAlaAsnAlaGlnLysTrpAlaAsnGlnCysAsnTrpArgHisSe 98
DB 411 GAGCAGAGAGGTAAACAGCAATGCCCAAGGTGGCAACAACTACTTACAA-CATAG 469
OY 98 rAsnProLysAspArgMetThrSerLeuLysCysGlyGluAsnLeuTrpMetSerSerAl 118
DB 470 TGATCCAGAGGAGCCGAAACAGTACAAAGATGCTGAGAAATCTATATGTCAGATGA 529
OY 118 aProSerSerTrpSerGlnAlaAlaGlnInsertPrpPheAspGluTrpAsnAspPheAspH 138
DB 530 CCTACTCTCTGCTCTTCTGCAATCCAAAGCTGGTATGACGAGATCCTAGATTTTGTCTA 589
OY 138 eGlyValGlyProLysThrProAsnAlaValAlaGlyHisLysTrpGlnValAlaTripty 158
DB 590 TGGTGTAGAGACCAAGAGTCCAA----- 613
OY 158 rSerSerTrpLeuValGlyCysGlyAsnAlaTrpCysProAsnGlnLysValLeuLysTrp 178
DB 613 ----- 613
OY 178 rTyTrpValCysGlnTrpCysProAlaGlyAsnTrpAlaAsnArgLeuTrpValProTy 198
DB 614 -----TATGTCTCTGCTGTATATATATGAAATAGAAAGATATACCCCGTA 657
OY 198 rGlnGlnGlyAlaProCys 204
DB 658 CCACAGAGGACACCTTGT 676
```

Search completed: March 14, 2003, 03:17:52
Job time : 393.449 secs